§ 452.115g

milliliters of methyl alcohol. Add 300 milliliters of 0.1M potassium phosphate buffer, pH 8.0 (solution 3), and blend again for 2 to 3 minutes. Hydrolyze this solution in a 60° C. constant temperature water bath for 2 hours or at room temperature for 16 to 18 hours. Further dilute with solution 3 to the reference concentration of 1.0 microgram of erythromycin base per milliliter (estimated).

(2) *Moisture.* Proceed as directed in §436.201 of this chapter.

[39 FR 19149, May 30, 1974, as amended at 50 FR 19921, May 13, 1985]

§452.115g Erythromycin estolate and sulfisoxazole acetyl oral suspension.

- (a) Requirements for certification—(1) Standards of identity, strength, quality, and purity. Erythromycin estolate and sulfisoxazole acetyl oral suspension is erythromycin estolate sulfisoxazole acetyl with suitable and harmless buffer substances, preservatives, solvents, stabilizers, emulsifiers, dispersing agents, colorings, flavorings. Each milliliter contains erythromycin estolate equivalent to 25 milligrams of erythromycin and 120 milligrams of sulfisoxazole. Its erythromycin content is satisfactory if it is not less than 90 percent and not more than 120 percent of the number of milligrams of erythromycin that it is represented to contain. Its sulfisoxazole acetyl content is satisfactory if it is not less than 90 percent and not more than 115 percent of the number of milligrams of sulfisoxazole that it is represented to contain. Its pH is not less than 3.5 and not more than 6.5. The erythromycin estolate used conforms prescribed the standards §452.15(a)(1). The sulfisoxazole acetyl used conforms to the standards prescribed by the U.S.P. XXII.
- (2) Labeling. It shall be labeled in accordance with the requirements of §432.5 of this chapter.
- (3) Requests for certification; samples. In addition to complying with the requirements of §431.1 of this chapter, each such request shall contain:
 - (i) Results of tests and assays on:
- (A) The erythromycin estolate used in making the batch for potency, moisture, pH, crystallinity, and identity.

- (B) The sulfisoxazole acetyl used in making the batch for all U.S.P. XXII specifications.
- (C) The batch for erythromycin content, sulfisoxazole content, and pH.
- (ii) Samples, if required by the Center for Drug Evaluation and Research:
- (A) The erythromycin estolate used in making the batch: 10 packages, each containing not less than 500 milligrams.
- (B) The batch: a minimum of 15 immediate containers.
- (b) Tests and methods of assay—(1) Erythromycin content. Proceed as directed in §436.105 of this chapter, preparing the sample for assay as follows: Remove an accurately measured representative volume of the suspension and dilute with sufficient methyl alcohol to give a concentration of 2.5 milligrams per milliliter (estimated). Dilute the entire mixture with sufficient 0.1M potassium phosphate buffer, pH 8.0 (solution 3), to give a concentration of 1.0 milligram of erythromycin base per milliliter (estimated). Hydrolyze in a 60 °C constant temperature water bath for 2 hours or at room temperature for 16 to 18 hours. Further dilute with solution 3 to the reference concentration of 1.0 microgram of erythromycin base per milliliter (estimated).
- (2) Sulfisoxazole content. Proceed as directed in § 436.328 of this chapter.
- (3) *pH*. Proceed as directed in §436.202 of this chapter, using the drug as it is prepared for dispensing.

[55 FR 280, Jan. 4, 1990]

§ 452.125 Erythromycin ethylsuccinate oral dosage forms.

§ 452.125a Erythromycin ethylsuccinate chewable tablets.

(a) Requirements for certification—(1) Standards of identity, strength, quality, purity. Erythromycin ethylsuccinate chewable tablets are composed of erythromycin ethylsuccinate and suitable and harmless diluents, binders, buffers, colorings, and flavorings. Each tablet conerythromycin ethylsuccinate tains equivalent to 200 milligrams of erythromycin. Its potency is satisfactory if it is not less than 90 percent and not more than 120 percent of the number of milligrams of erythromycin that it is